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The effects of croscarmellose sodium concentration on the physicochemical characteristics of orodispersible tablets of atenolol**Nani Parfati, Karina Citra Rani****Department of Pharmaceutics, Faculty of Pharmacy,**University of Surabaya, Surabaya**Jl. Kalirungkut Surabaya, Surabaya-60293, Jawa Timur, Indonesia**Submitted: 18-10-2017**Reviewed: 20-03-2018**Accepted: 29-04-2018***ABSTRACT**

Hypertension is the most common cardiovascular diseases suffered by geriatric patients. Their physiological changes make the administration of conventional tablets less effective, especially regarding compliance. One approach to overcome this problem is the development of orodispersible tablets, which soften easily and disintegrate quickly in the oral cavity. Atenolol is a class of β -blocker functioning as an anti-hypertensive drug that has been extensively used in hypertension therapy, and it has the potential to being developed as orodispersible tablets. A faster disintegration of orodispersible tablets will facilitate an earlier onset of dissolution. The addition of superdisintegrants can reduce the disintegration time of these tablets. Croscarmellose sodium is a superdisintegrant that can decrease the disintegration time to less than three minutes. This study aimed to optimize the formula of orodispersible tablets of atenolol using different concentrations of croscarmellose sodium, namely 10% (formula 1) and 20% (formula 2). The physicochemical characteristics of the tablets were evaluated to determine the best formula. The evaluation included a comparison to the control formula (0% of croscarmellose sodium). The results showed that formula 1 (10% of croscarmellose sodium) produced orodispersible tablets with the best physicochemical characteristics regarding tablet hardness, friability, in vitro dispersion time, and disintegration time. Formula 1 (% $Q_{30 \text{ minutes}}$ = 98.31%) also met the standard of the dissolution of atenolol tablets set by the Farmakope Indonesia, i.e., the percent of dissolved drug in 30 minutes has to be higher than 85%. The drug dissolution efficiency of formula 1 was twice higher than that of the control formula.

Keywords: atenolol, croscarmellose sodium, hypertension, orodispersible tablets

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INTRODUCTION

Hypertension is one of the degenerative diseases suffered by geriatric patients. It is a cardiovascular disease with increased prevalence in older adults (Dipiro *et al.*, 2008). Atenolol is a β -blocker (β 1-selective) antihypertensive drug that is widely used for the treatment of all lines of hypertension, angina pectoris, arrhythmias, and myocardial infarction (Dipiro *et al.*, 2008). It is also an antihypertensive drug listed in the formulary of the Social Security Administering Agency (BPJS Indonesia) of Health. In oral administration, it is absorbed by approximately 50%, and most of the absorbed doses reach the systemic circulation. The half-life of atenolol is 6-7 hours with t_{\max} between 2-4 hours (Sweetman, 2009). Atenolol is a compound with low water solubility and dissolution but high permeability (Khirwadkar and Dashora, 2013). The low water solubility results in a slow dissolution process that affects the onset of drug action and atenolol bioavailability in the systemic circulation. The conventional administration of atenolol tablets with a glass of water causes discomfort to geriatric patients as they experience physiological changes, such as difficulty in swallowing and chewing, tremor, and risk of choking (Khirwadkar and Dashora, 2013).

The development of atenolol as orodispersible tablets is an approach to reduce the time required for tablet disintegration and minimize the problems in the conventional administration of tablets, especially in geriatric patients. Orodispersible tablet or orally disintegrating tablet (ODT) is a dosage form that quickly dissolves into fine particles when placed in the oral cavity (European Pharmacopoeia Commission, 2005). It can be absorbed in the mouth, pharynx, and esophagus and dissolved and dispersed in saliva in less than three minutes (European Pharmacopoeia Commission, 2005). Furthermore, ODT has several advantages, including fast dissolution process and drug action, minimum first-pass effect, and increased drug bioavailability (Chandrasekhar *et al.*, 2013). It also facilitates patients in taking medicine and, thereby, increases their drug compliance because (1) it is easily swallowed and effective, (2) it has a relatively good flavor, and (3) patients do not need to chew and drink it with water (Kumare *et al.*, 2013). Therefore, this research developed the formulation of atenolol for orodispersible tablets.

An essential component in the formulation of orodispersible tablets as a dosage form is disintegrant. Selecting a suitable disintegrant is, therefore, necessary mainly to create ODT formulas that meet the required physicochemical characteristics. The use and the selection of disintegrants have to be precise in order to obtain the desired dissolution and disintegration rates (Sulaiman, 2007; Camarco *et al.*, 2006). Disintegrant is one of the additional ingredients in drug formulation that serves to break the tablet into smaller particles, enabling fast dissolution process. A group of disintegrant used in low level in the preparation of tablets is termed superdisintegrant. It is considered to be effective as an excipient in ODTs when applied with concentrations between 10% and 20% (Hahm and Augsburg, 2008). One type of superdisintegrant that can be used in ODT formulation is croscarmellose sodium. (Hahm and Augsburg, 2008). Croscarmellose sodium facilitates the disintegration process through the mechanism of swelling, recovery of elastic energy, and capillary action (wicking). It is a fiber-like polymer with a rather short size and adequate flow characteristics. Its long fiber-shaped structure can widen the distance between the constitutive particles of the matrix tablet, which accelerates the disintegration process (Hahm and Augsburg, 2008). Accordingly, a thorough study related to the effects of croscarmellose sodium concentration in the ODT formula on the physicochemical characteristics of the produced ODT.

This research formulated orodispersible tablets of atenolol with croscarmellose sodium as superdisintegrant at different concentrations, namely 10% (formula 1) and 20% (formula 2). It also used a control formula, i.e., an orodispersible tablet of atenolol prepared without the addition of croscarmellose sodium (0%). The formulation stage included pre-compression and post-compression evaluations. The results of this stage determined the most favorable formula based on the physical characteristics of ODT and the dissolution of atenolol. The addition of different concentrations of croscarmellose sodium as a disintegrant was analyzed for its impact on the physicochemical characteristics of ODT, i.e., disintegration time, dispersion time, tablet hardness, and atenolol level.

Furthermore, the pharmaceutically qualified formulas with the best physical properties were tested for their dissolution rate according to the standards issued in the Farmakope Indonesia V (Depkes RI, 2014).

MATERIALS AND METHODS

Materials

The materials used in this study included pharmaceutical components, excipients, and reagents. All of the components and excipients were pharmaceutical grade. The components were atenolol, mannitol DC (direct compress), croscarmellose sodium, Avicel PH 102®, magnesium stearate, Aerosil®, aspartame, and talc. The reagents and other chemicals included methanol (Merck), sodium dihydrogen phosphate (Merck), disodium hydrogen phosphate (Merck), sodium acetate (Merck), and glacial acetic acid (Merck). All of the reagents and chemicals were analytical grades.

Preparation of the pressed mass

The composition of the ingredients used in the pressed mass is presented in Table I. This study used three formulas for the production of orodispersible tablets of atenolol with croscarmellose sodium as superdisintegrant. The formulas included a control (0%), formula 1 (10% of croscarmellose sodium), and formula 2 (20% of croscarmellose sodium). The pressed mass of each formula was prepared by initially mixing atenolol with some of Aerosil® for 3 minutes and then with some of the Avicel PH 102® for 5 minutes. The next step was the addition of croscarmellose sodium as the superdisintegrant with different concentrations according to the formulas (0%, 10%, or 20%), aspartame, mint flavor, the rest of the Avicel PH 102®, and mannitol DC. These components were mixed in a tumbling mixer until homogeneous for 10 minutes. This process produced the powder form of the orodispersible tablets before compression.

Table I. The formulas of the orodispersible tablets of atenolol with croscarmellose sodium at concentrations of 0% (control), 10% (formula 1), and 20% (formula 2)

Ingredients	Control Formula (mg)	Formula 1 (mg)	Formula 2 (mg)
Atenolol	25	25	25
Croscarmellose sodium	-	30	60
Aspartame	9	9	9
Mg Stearate	4.5	4.5	4.5
Aerosil	1.5	1.5	1.5
Talc	3	3	3
Mint Flavor	3	3	3
Mannitol DC	50.8	44.8	38.8
Avicel PH 102®	203.2	179.2	155.2
Total weight per tablet	300	300	300

Pre-compression evaluation

After the powder form of the orodispersible tablets was created, a pre-compression evaluation was performed. It analyzed the flow rate, angle of repose, compressibility index, Hausner ratio, and moisture content of the powder.

Flow rate and angle of repose

The flow rate and angle of repose were identified by weighing 100 grams of the powder form. This powder was poured into a funnel set on a stative, and the distance between the lower end of the

pipe to the flat plane was 10.0 ± 0.2 cm. The lower hole of the funnel was closed. This hole was then opened later, and the time required for the material to flow out of the funnel was recorded with a stopwatch for flow rate calculation. The next step was measuring the height of the material pile and the radius of the cone base, as well as calculating the resultant angle of repose (Aulton and Summers, 2013).

Compressibility index

The compressibility index of the powder form was determined by measuring two types of bulk density, namely freely settled bulk density and tapped bulk density of the powder form (Aulton and Summers, 2013). Both data were used to calculate the compressibility index of the powder using the following equation:

$$\text{Compressibility Index} = \frac{\rho_T - \rho_B}{\rho_T} \times 100\%$$

where:

ρ_T : freely settled bulk density of the powder

ρ_B : tapped bulk density of the powder

Hausner ratio

Hausner ratio compares the tapped bulk density with the freely settled bulk density of the powder (Aulton, 2013). Both Hausner ratio and compressibility index were used to predict the flowability of the powder.

Moisture content

The moisture content was measured by weighing 5 grams of the powder form. The surface of the powder was flattened on the container. The position of the heating lamp was arranged in a way that it was above the material. During the drying process, the weight of the powder was displayed every 15 minutes.

A drying process is deemed perfect when the weight of the powder does not change for 3x15 minutes (Aulton & Summers, 2013). Moisture content was calculated using the following formula:

$$\%MC = \frac{W - W_0}{W_0} \times 100\%$$

where:

MC : moisture content

W : wet weight of the powder

W_0 : dry weight of the powder

Compression of orodispersible tablets of atenolol

Before the compression process from a powder into a tablet dosage form, the powder was mixed first with the external phase, i.e., magnesium stearate, talc, and the remaining Aerosil® using a tumbling mixer for three minutes until homogeneous. This powder mixture was then compacted into a tablet dose using ERWEKA® AR 402 tablet compression machine, and the weight of each tablet was 300 mg. The diameter of the punch was 11 mm, the surface of the punch was round and biconvex, and the compression force was 2 tons.

Post-compression evaluation

After the powder form was pressed into a tablet dose, a post-compression assessment was performed. Aside from organoleptic analysis, the evaluated properties were atenolol content, hardness, disintegration time, tablet dispersion time, friability, and dissolution.

Organoleptic analysis

This analysis was conducted by observing the physical appearance of every tablet produced by each formula, including the odor, color, flavor, and shape of the compressed orodispersible tablets of atenolol.

Identification of atenolol content

The atenolol levels were assessed by sampling 20 tablets of each formula randomly. The twenty samples were crushed together. A mixture of atenolol that was equivalent to 25 mg was weighed and dissolved with 10 mL of methanol in a measuring flask, then added with acetate buffer solution (pH 4.6) up to 100 mL. Ten mL of this solution was transferred with a pipet into a measuring flask and added with acetate buffer solution up to 25 mL. This solution was filtered with Whatman paper No.41. The absorbance of the filtered solution was measured with UV-VIS spectrophotometer at a wavelength of 274 nm using an acetate buffer blank (pH 4.6) (Chandrasekhar *et al.*, 2013).

Evaluation of tablet hardness

The tablet hardness was evaluated using the instrument 'Monsanto hardness tester'. The number of samples collected from each formula for this evaluation was ten tablets.

Disintegration time test

Disintegration time is a critical parameter to observe in the evaluation of orodispersible tablets. The disintegration time was tested by placing 6 tablets (of each formula) into 6 different tubes containing 900 mL of water with a temperature of $37 \pm 0.5^\circ\text{C}$. As the test apparatus started, the stopwatch was turned on. The length of time that the tablet required to disintegrate, or the duration that the core mass needed to become palpable, was determined as the disintegration time of the tablet (Chandrasekhar *et al.*, 2013; Kumare *et al.*, 2013).

In vitro dispersion time test

Each tablet from the three formulas was placed in a beaker containing 6 mL of phosphate buffer solution (pH 6.8) with a temperature of $37 \pm 0.5^\circ\text{C}$. The time required by the tablet to crumble was recorded as the in vitro dispersion time (Chandrasekhar *et al.*, 2013; Kumare *et al.*, 2013).

Friability and abrasion tests

The friability and abrasion tests were performed by selecting several tablets until a weight of 6.5 gram was reached, i.e., approximately 22 tablets (for tablets weighing lower than 650 mg). These tablets were weighed using an analytical balance. Afterward, they were placed into the friability and abrasion tests instrument, which was operated at 25 rpm for 4 minutes. The weight of the entire tablets was recorded, and the weight difference between the tablets before and after the test was calculated (Chandrasekhar *et al.*, 2013; Aulton and Summers, 2013). The percentage of friability or abrasion was calculated using the following formula:

$$\% \text{friability} = \frac{\text{Initial weight} - \text{Final weight}}{\text{Initial weight}} \times 100\%$$

Dissolution test

The dissolution test of the orodispersible tablets of atenolol was performed on a dissolution media, i.e., 900 mL of 0.1 N acetic acid buffer (pH 4.6), according to the standardized procedure issued in Farmakope Indonesia V (Depkes RI, 2014). The dissolution test apparatus used the type 2 (paddle) at 50 rpm. The first step was inserting six tablets of each formula into the apparatus. Then, 10 mL of samples was drawn every time of observation, namely 0, 1, 2, 3, 4, 5, 10, 15, 20, 25, 30, 45, 60,

75, 90, 105, and 120 min. Afterward, 10 mL of 0.1 N acetic acid (pH 4.)⁶ was returned to the apparatus. The next step was assessing the atenolol level of each sample using a spectrophotometer at a wavelength of 274 nm. The dissolution parameter (% Q, tQ%, the area under the dissolution curve (AUC), and dissolution efficiency) and dissolution profile were identified based on the readings (Chandrasekhar *et al.*, 2013). The dissolution of AUC was determined by calculating the area formed under the dissolution profile. Dissolution efficiency was defined as the area under the dissolution curve until a certain time limit and expressed as a percentage of the rectangle are depicted by 100% dissolution at the same time limit (Khan, 1975).

Data Analysis

The post-compression evaluation results of the orodispersible tablets of atenolol created with the control formula (0% of croscarmellose sodium), formula 1 (10% of croscarmellose sodium), and formula 2 (20% of croscarmellose sodium) were statistically analyzed using One-way ANOVA with 95% confidence level ($\alpha = 0.05$). The examined parameters included disintegration time and in vitro dispersion time. The AUC dissolution and dissolution efficiency of the selected formula were compared with the control formula using a pooled t-test.

RESULTS AND DISCUSSION

The orodispersible tablets of atenolol in this research were prepared using croscarmellose sodium as superdisintegrant. The formula designed in this study aimed to determine the effects of different concentrations of croscarmellose sodium on the physicochemical characteristics of the resulted tablets. A control formula was designed without superdisintegrant. Meanwhile, formula 1 and formula 2 used 10% and 20% of croscarmellose sodium, respectively. The pre-compression evaluation was performed on the powder form of these three formulas. It included flow rate, angle of repose, compressibility index, Hausner ratio, and moisture content. The results of the pre-compression evaluation are presented in Table II.

Table II. The results of the pre-compression evaluation of the orodispersible tablets of atenolol with different concentrations of croscarmellose sodium (0%, 10%, and 20%)

Parameters	Formulas*			Standards
	Control (0%)	Formula 1 (10%)	Formula 2 (20%)	
Flow rate (g/s)	9.31±0.16	7.10±0.82	7.38±0.31	4-10
Angle of repose (°)	32.15±0.00	27.84±01.35	29.53±0.29	5°-35°
Compressibility (%)	23.69±0.499	25.48±1.07	27.55±0.36	< 25%
Hausner Ratio	1.32±0.006	1.34±0.02	1.38±0.01	< 1.25
Moisture content (%)	4.80±0.234	5.11±0.15	5.28±0.56	3-5%

*The figures represent the average values

The flow rates of the three formulas were between the range of 4-10 g/s, indicating sufficient flow characteristics (Hahm and Augsburg, 2008; Aulton and Summers, 2013). The angle of repose was employed in this study to characterize the flow of a solid particle as it served as an indicator of interparticle friction or movement resistance between particles (United States Pharmacopeia Convention, 2017). Based on the angle of repose, both formula 1 and formula 2 had an excellent flow characteristic, whereas the control formula was categorized as a powder with sufficient flow.

The assessment of compressibility index and Hausner ratio aimed to identify the characteristics of the powder form during tableting. The threshold for good powder compressibility is smaller than 20% (Agoes, 2012). The Hausner ratio can be divided into three groups, namely excellent flow (1.00-1.11), good flow (1.12-1.18), and sufficient flow (1.19-1.25). The results of both compressibility index

and Hausner ratio categorized the control formula and formula 1 as a passable powder or, in other words, a powder could flow. On the contrary, formula 2 was concluded as a category of powder with poor flow. Based on the angle of repose, Hausner Ratio, and compressibility index, the powder form had a good flow with some obstacles. The poor Hausner ratio and compressibility index were caused by the presence of a large number of fine particles in the powder (Aulton and Summers, 2013). The moisture contents of control formula, formula 1, and formula 2 were 4.80%, 5.11%, and 5.28%, respectively. The presence of hygroscopic excipient caused such high moisture content. The superdisintegrant used in this research had a high affinity for water so that the powder absorbed moisture from the air (Aulton and Summers, 2013). The moisture content of a powder form has to be controlled to prevent sticking or picking on the punch during the tablet compression process.

The post-compression evaluation in this study included organoleptic examination and tests of atenolol level, tablet hardness, disintegration time, in vitro dispersion time, friability, abrasion, and dissolution. The results are summarized in Table III. The organoleptic characteristics of control formula, formula 1, and formula 2 were round, white, sweet flavor, and minty odor. The physical appearance of the orodispersible tablets of atenolol produced in this study is presented in Figure I.

The atenolol contents in control formula, formula 1, and formula 2 met the thresholds set by the Farmakope Indonesia, namely 90%-110% (Depkes RI, 2014). The orodispersible tablets of atenolol must have sufficient physical strength to withstand various mechanical shocks during the manufacturing, packaging, and delivery processes. The orodispersible tablets of atenolol produced in this research had an excellent hardness between the range of 2-4 kg (Deshmukh *et al.*, 2012; Sharma, 2013). The three formulas met the standards for ODT hardness.



Figure 1. The orodispersible tablet of atenolol with croscarmellose sodium as superdisintegrant

Table III. The results of the post-compression evaluation of the orodispersible tablets of atenolol with different concentrations of croscarmellose sodium (0%, 10%, and 20%)

Parameters	Formulas*			Standards
	Control (0%)	Formula 1 (10%)	Formula 2 (20%)	
Organoleptic	Round, white, sweet flavor, minty odor	Round, white, sweet flavor, minty odor	Round, white, sweet flavor, minty odor	Round, white, sweet flavor, minty odor
Atenolol level (%)	93.99±0.13	109.10±0.13	108.51±0.22	90.0-110.0%
Hardness (kP)	2.85±0.24	3.15±0.58	2.90±0.21	2-4 Kp
Disintegration time (s)	4.00±0.00	2.57±0.00	2.89±0.00	< 3 minutes
In vitro dispersion time	15.00±0.00	4.44±0.41	7.29±0.56	-
Friability (%)	0.20±0.09	0.07±0.03	0.17±0.08	< 1%
Abrasion (%)	0.25±0.18	0.07±0.02	0.07±0.02	< 1%

*The figures represent the average values

This research also evaluated the disintegration time and the in vitro dispersion time of the produced tablets. Favorable disintegration time for orodispersible tablets of atenolol is less than 3 minutes (≤ 180 seconds) (European Pharmacopoeia Commission, 2005). All of the three formulas met the standards of tablet disintegration time in the purified water. Formula 1, containing 10% of croscarmellose sodium, had the shortest disintegration time. In vitro dispersion time predicts the ability of an orodispersible tablet to break into fine particles with a very limited volume of saliva (± 6 ml). The statistical analysis, i.e., One-way ANOVA, showed that there was a significant difference of disintegration time between the three formulas ($p < 0.05$). Furthermore, the results of the post hoc Tukey test showed that the dispersion time of formula 1 (10% of croscarmellose sodium) was faster than the other formulas. A rapid dispersion time indicates that the tablet is immediately dispersed into fine particles when in contact with saliva in the oral cavity (Nagendrakumar *et al.*, 2010). Based on the evaluation of the physical characteristics, formula 1 (10% of croscarmellose sodium) produced relatively fast disintegration time compared to control formula and formula 2. Increased concentration of croscarmellose sodium from 10% to 20% caused both disintegration time and in vitro dispersion time become longer because this superdisintegrant expanded when in contact with aqueous media. This condition triggers the formation of viscous barrier on the tablet surface, inhibiting the penetration of water into the tablet (Desai *et al.*, 2014).

Based on the physical properties, formula 1 (10% of croscarmellose sodium) is the best formula regarding pre-compression and post-compression characteristics (disintegration time and in vitro dispersion time). Therefore, the dissolution properties of formula 1 were identified, and the results were compared with the dissolution of the control formula. The dissolution profiles of formula 1 and the control formula are depicted in Figure 2, while the calculation results of the dissolution parameters of both formulas are summarized in Table IV.

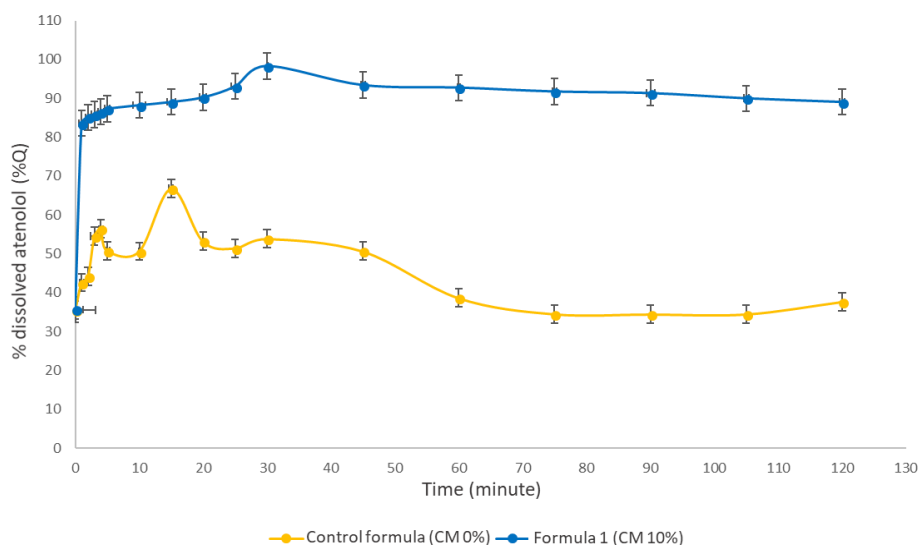


Figure 2. The dissolution profiles of orodispersible tablets of atenolol produced with control formula (0% of croscarmellose sodium) and formula 1 (10% of croscarmellose sodium)

Table IV. The dissolution parameters of the orodispersible tablets of atenolol produced with the control formula and formula 1

Dissolution Parameters	Control Formula	Formula 1
AUC	5,190.32±2.78	10,964.99±27.99
Dissolution efficiency (ED _{120 minutes} (%))	43.25±0.02	91.37±0.23
%Q (30 minutes)	53.85	98.31%
tQ% (minutes)	270.84	1.91
K dissolution	0.1893/minute	0.4150/minute

Based on the unpaired t-test, the AUC values and dissolution efficiencies of formula 1 and the control formula are different ($p < 0.05$). The dissolution efficiency of formula 1 (10% of croscarmellose sodium) is twice higher than the control formula. Furthermore, the orodispersible tablets of atenolol prepared using formula 1 met the requirements of ODT dissolution. The amount of dissolved atenolol (formula 1) in 30 minutes was 98.31%, which is higher than the standard set by the Farmakope Indonesia, i.e., 85% (Depkes RI, 2014).

CONCLUSIONS

In this study, the different concentrations of croscarmellose sodium as superdisintegrants in the formulation of orodispersible tablets of atenolol affect the physicochemical properties of the ODT. Orodispersible tablets of atenolol prepared with 10% of croscarmellose sodium (formula 1) exhibit optimum results in both pre-compression and post-compression evaluations. This formula successfully produces ODT of atenolol with the fastest disintegration time and the highest dissolution compared to the other formulas.

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